

GB1209326

Publication Title:

THE PREPARATION OF 3-METHYL-5-PYRAZOLE CARBOXYLIC ACID AMIDES

Abstract:

Abstract of GB1209326

1,209,326. 3-Methyl-5-pyrazolecarboxylic acid amides. CARTER WALLACE Inc. 21 Sept., 1967 [27 Sept., 1966], No. 42957/67. Heading C2C. [Also in Division A5] 3 - Methyl - 5 - pyrazolecarboxylic acid amides of the formula wherein R is NH₂ or a radical of a substituted o<SP>r</SP> unsubstituted primary or secondary amine (the compounds in which R in the above formula represents morpholino, piperidono, benzylamino and p-chlorobenzylamino being novel) are prepared by treating ethyl acetopyruvate, dissolved in glacial acetic acid, with hydrazine hydrate, cooling to cause ethyl 3-methyl-5- pyrazole carboxylate to precipitate, reacting said ester with ammonia or primary or secondary amine in stoichiometrically equivalent proportions in an aqueous alcoholic solution, in an excess of the amine, or in aqueous solution containing an excess of the amine or ammonia, freuxing for 4 to 15 hours and collecting the precipitated amide. Reference has been directed by the Comptroller to Specification 1,048,104. Data supplied from the esp@cenet database - Worldwide

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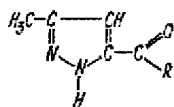


(54) THE PREPARATION OF 3-METHYL-5-PYRAZOLE
 CARBOXYLIC ACID AMIDES

(71) We, CARTER WALLACE, INC., a corporation organized and existing under the laws of the State of Maryland, United States of America of Two Park Avenue, New York, N.Y.10016, United States of America, do hereby declare the invention, for which we pray that a patent may be granted, to us, and the method by which it is to be performed, to be particularly described in and by the following statement:—

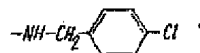
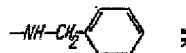
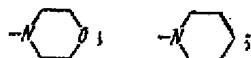
This invention relates to the preparation of 3 - methyl - 5 - pyrazolecarboxylic acid amides.

The products produced by the process of the invention have the following general formula:



wherein R is NH₂ or a radical of a primary or secondary amine, such as

20



These compounds may be used in association with a pharmaceutical carrier in pharmaceutical compositions for the purposes described below. 25

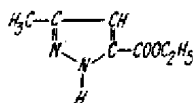
The amides of formula I are prepared according to the invention by a process comprising the steps of treating ethyl aceto-pyruvate, dissolved in glacial acetic acid, with hydrazine hydrate, cooling to cause 3 - methyl - 5 - pyrazolecarboxylic acid ethyl ester to precipitate, reacting said ester with ammonia or a primary or secondary amine in stoichiometrically equivalent proportions in an aqueous alcoholic solution, in an excess of the amine, or in aqueous solution containing an excess of the amine or ammonia, refluxing for 4 to 15 hours and collecting the precipitated amide. After the refluxing the amide is precipitated either by concentration or cooling the solution. The precipitate is recrystallised two or three times, as necessary. 40

The ethyl ester of the 3 - methyl - 5 - pyrazolecarboxylic acid is a known compound, described by Knorr (Annalen, 279, page 219) of 1894 and the process of its preparation has been described in the German Patent No. 74,619. 45

This ester, however, when prepared by Knorr's method, that is the reaction of hydrazine sulphate with the sodium salt of ethyl aceto-pyruvate in the presence of NaOH in 50

aqueous solution, does not exhibit constant characteristics and so does not lend itself to the satisfactory preparation of the compounds of formula I.

- 5 The novel method for the preparation of the ethyl ester of 3 - methyl - 5 - pyrazolecarboxylic acid, having the formula:



- 10 used in this invention is, in more detail, as follows:

To 5 g. (0.032 mol) of ethyl aceto-pyruvate (prepared as disclosed in A. H. Blatt, "Organic Syntheses Coll. Vol. I, published by John Wiley & Sons, New York, 1958, page 233) dissolved in 5 mls. glacial acetic acid, 1.85 ml. of 85% hydrazine hydrate are added gradually and with stirring (0.032 mol.).

The solution is refluxed for 15—20 mins. and on cooling to room temperature, is poured into an ice-filled vessel. A flaky white precipitate is formed which is recrystallised from ligroin.

There are obtained 3.1 g. (yield 62%) of the ethyl ester 3 - methyl - 5 - pyrazolecarboxylic acid which is soluble in ligroin, poorly soluble in ethanol, diethyl ether and benzene, soluble in water only with difficulty. Melting point 81°C—82°C. (The chemical literature reports a m.p. of 82°—83°C.).

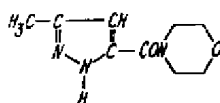
30 By changing either the primary or the secondary amine used in the preparation many compounds can be obtained which correspond to the general formula reported above.

65 The product analyses:

For $C_8H_{11}N_3O$ Calculated: C% = 47.99; H% = 5.64; N% = 33.58
Found: C% = 47.79; H% = 5.61; N% = 33.15

EXAMPLE 2

70 3 - methyl - 5 - pyrazolecarboxylic acid morpholide



75 5 g. (0.032 mol) of the ethyl ester of 3 - methyl - 5 - pyrazolecarboxylic acid are refluxed with an excess of morpholine (10 ml. approx.) for about 4 hours. The mixture is

Analysis:

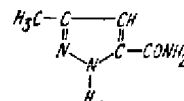
90 For $C_9H_{13}N_3O_2$ Calculated: C% = 55.37; H% = 6.71; N% = 21.52
Found: 55.67 6.67 21.43

Of the compounds of formula I the morpholide, piperidide, benzylamide, and *para*-chlorobenzylamide of 3 - methyl - 5 - pyrazolecarboxylic acid are novel and as such within the scope of the invention.

In the following Examples, a few of these compounds are described along with the procedure for their preparation, by way of example only and without any implied limitation thereto.

EXAMPLE 1

3 - methyl - 5 - pyrazolecarboxylic amide 45



6 g. (0.039 mol) of the ethyl ester of 3 - methyl - 5 - pyrazolecarboxylic acid are dissolved in the minimum possible amount of ethanol and then treated with an excess of aqueous ammonia.

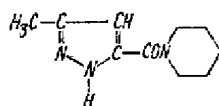
The mixture is refluxed for 15 hours and a white solid precipitates on cooling the solid which is twice recrystallised from water, has one molecule of water of crystallisation. (Analysis: Calculated for $C_8H_9N_3O \cdot H_2O$, N% = 29.34; Found: N% = 29.23).

The product when heated for 6 hours in an oven under reduced pressure, loses its molecule of water. Yield 3.5 g. (63%). The desired anhydrous product melts at 174°C. and is poorly soluble in cold water and ethanol while it is fairly soluble in the same solvents when hot.

diluted with water and then the water and morpholine are removed by heating under reduced pressure.

The residue thus obtained is recrystallised in the presence of decolorising carbon and dried in a vacuum oven. According to the best procedure, it is recrystallised twice from ethanol and precipitated with diethyl ether. There are obtained 3 g. (yield 50%) of the desired product as a white solid which melts at 188°C. It is soluble in water and ethanol, and insoluble in diethyl ether.

EXAMPLE 3

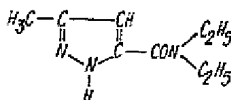
3 - methyl - 5 - pyrazolecarboxylic acid piperi-
dide

The same procedure as in Example 2 is followed except, instead of morpholine, an excess of piperidine (10 ml. approx.) is used. According to the best procedure, the product is twice recrystallised from aqueous ethanol. The desired product thus obtained (yield 53%) melts at 191°C—192°C. and is soluble in ethanol and poorly soluble in water.

The product analyses:

For $C_{10}H_{13}N_3O$ Calculated: C% = 62.15; H% = 7.82; N% = 21.75
Found: 62.05 7.63 23.58

EXAMPLE 4

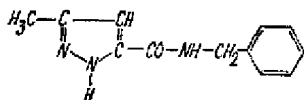
3 - methyl - 5 - pyrazolecarboxylic acid di-
ethylamide

The same procedure as in Example 2 is followed except, instead of morpholine, an excess of diethylamine (10 ml. approx.) is used. According to the best procedure, the desired product is recrystallised twice from alcohol and precipitated with ether. A white solid (yield 53%) which melts at 164°C. and is soluble in water and ethanol, and insoluble in diethyl ether is obtained.

The product analyses:

For $C_{10}H_{15}N_3O$ Calculated: C% = 59.64; H% = 8.34; N% = 23.18
Found: 59.36 8.12 23.21

EXAMPLE 5

3 - methyl - 5 - pyrazolecarboxylic acid
benzylamide

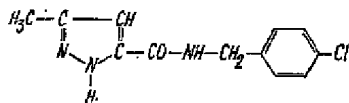
5 g. (0.032 mol.) of the ethyl ester of 3 - methyl - 5 - pyrazolecarboxylic acid, dissolved in aqueous ethanol, are reacted with 3.3 g. of benzylamine by refluxing for 5 hours.

The solution is evaporated and the desired product thus obtained, when recrystallised twice from aqueous ethanol, melts at 116°C. and is soluble in ethanol, and insoluble in water. Yield 63%.

The product analyses:

For $C_{12}H_{13}N_3O$ Calculated: C% = 66.95; H% = 6.09; N% = 19.52
Found: 67.42 6.06 19.67

EXAMPLE 6

3 - methyl - 5 - pyrazolecarboxylic acid - p -
chlorobenzylamide

The procedure of Example 5 is followed by employing 4.3 g. of p-chlorobenzylamine and the precipitate thus obtained upon cooling is recrystallised twice from aqueous ethanol. The desired product melts at 202°C. and is poorly soluble in ethanol and insoluble in water. Yield 63%.

The product analyses:

For $C_{12}H_{12}ClN_3O$ Calculated: C% = 57.72; H% = 4.64; N% = 16.83
Found: 58.20 4.69 16.61

All the compounds obtained according to the Examples from 1 to 6 are endowed with outstanding pharmacological properties such as an activity upon ketone bodies, lipids and triglycerides.

In the following the results of a pharmaco-

logical screening, aiming at ascertaining the activity of the 3 - methyl - 5 - pyrazolecarboxylic acid amide ($CONH_2$) are reported in comparison with 3,5 - dimethyl pyrazole (3.5 DMP).

Activity of 3 - methyl - 5 - pyrazole-carboxylic acid amide on the lipid metabolism.

Methods — Male Sprague-Dawley rats, 150 g. average weight, were used. In "in vivo" experiments 3,5-dimethylpyrazole (3.5 DMP) and 3 - methyl - 5 - pyrazolecarboxylic acid amide (CONH₂) were given to 18 hours fasted rats orally or intraperitoneally.

At the end of the experiment the levels of plasma free fatty acids (FFA), ketone bodies, triglycerides (TG), cholesterol (CHOL), phospholipids (P), of blood glucose and of liver triglycerides were determined.

FFA were determined according to Dole (J. Clin. Invest. 1956, 35, 150) with minor modification. Lipids were extracted with a mixture of chloroform and methanol (2:1) and washed with saline. Phosphorus of phospholipids was determined in the chloroform extract according to Lowry *et al* (J. Biol. Chem. 1954, 207, 1). The residual chloroform extracts were shaken with silicic acid and centrifuged. Cholesterol and triglycerides were determined in the supernatant liquid according to the Lieberman and Burchard reaction and to the Van Handel and Zilversmit method (J. Lab. Clin. Med. 1957, 50, 153) respectively.

A further experiment was performed, during which rats were given CONH₂ each day for 15 days.

In other experiments CONH₂ and 3.5 DMP were given to rats treated with ethyl alcohol (1.6 ml./kg. *per os*).

Results — Table I gives the effect of CONH₂ on plasma FFA and ketone bodies. 18 hours fasted rats were given CONH₂ orally or intraperitoneally. Determinations were performed 30 min. or 60 min. after administration. CONH₂, given orally or intraperitoneally is effective in lowering plasma FFA and ketone bodies.

Table 2 gives the effect of CONH₂ and 3.5 DMP on plasma lipids, blood glucose and

liver TG. The drugs were administered by oral route to 18 hour fasted rats and the determinations were performed 6 hr and 8 hours after the treatment. 6 hours after the administration the lowering effect of 3.5 DMP on plasma lipid is already ended while CONH₂ is still active. 8 hours after CONH₂ administration the plasma FFA levels rose again, but plasma and liver TG were still low.

The CONH₂ does not show any effect on blood cholesterol or phospholipids. Results in Tables 3 and 4 are concerned with a longer course of administration of CONH₂; the rats were divided in 2 groups. The first group received CONH₂ 7.5 mg./kg. by oral route every day for 15 days. The second group received saline. At the end of the treatment each group was further divided into 3 groups: the first one was fed, the second was fasted for 18 hours, the third received CONH₂ 1 hour before being killed. Results in Table 3 show that the growth rate of CONH₂ treated rats was the same as that of control rats, and also that the adipose tissue weight was not affected. Results in Table 4 show that the effect of a shorter course of administration of CONH₂ was not affected by the previous longer treatment.

Results in Table 5 show that CONH₂ as well as 3.5 DMP prevented the increase of liver TG induced by ethyl alcohol.

In Table 6 it is shown that only CONH₂, but not 3.5 DMP, is curative with respect to a fatty liver induced by alcohol.

In Summary — CONH₂ decreases FFA probably by blocking lipolysis in the adipose tissue. CONH₂ decreases plasma and liver TG and in addition it lowers the level of plasma, and ketone bodies. CONH₂ prevents and cures the fatty liver induced by a single dose of ethyl alcohol. 3.5 DMP, the reference drug, prevents but it does not cure this toxic effect of alcohol.

TABLE 1

Effect of CONH_2 on plasma FFA and ketone bodies on fasted rats

Treatment mg/kg	route	Time between treatment and killing	Plasma	
			FFA $\mu\text{Eq/l}$	Ketone bodies mg/100 ml($^{\circ\circ}$)
Saline	—	—	699 ± 18	14.0 ± 0.5
CONH_2 1	i.p.	30'	235 ± 24	8.5 ± 0.6
Saline	—	—	554($^{\circ}$)	10.1 ± 0.9
CONH_2 7.5	os	60'	143	4.2 ± 0.5
3.7	"	"	143	3.6 ± 0.3
1.7	"	"	172	4.2 ± 0.1
0.75	"	"	168	4.4 ± 0.2
Saline	—	—	616($^{\circ}$)	$15. - \pm 2. -$
CONH_2 0.75	os	60'	265	6.3 ± 0.7
0.37	"	"	386	5.8 ± 0.6

($^{\circ\circ}$) determinations were performed on pooled sera.($^{\circ}$) each figure is the average of at least 5 determinations.

TABLE 2
Effect of CONH_2 and 3.5 DMP on plasma lipids blood glucose and Liver TG.

Treatment mg/kg, oral	time between treatment and killing	PLASMA					Blood glucose mg/100 ml	Liver TG mg/100 ml
		FFA	TG	CHOL	P			
		uEq/l	mg/100 ml	mg/100 ml	mg/100 ml			
Saline	—	602±37	73±6	57±5	3.4±0.2	55±3	695±56	
CONH ₂	7.5	345±38	32±1	60±4	3.3±0.2	56±4	433±15	
3.5 DMP	7.5	746±24	75±7	68±7	4.1±0.2	41±1	774±26	
Saline	—	872±74	69±4			63±2	408±25	
CONH ₂	7.5	656±64	46±2			69±3	288±20	
Saline	—	759±56	63±4			50±2	492±35	
CONH ₂	7.5	403±34	32±2			54±2	363±28	
Saline	—	897±64	74±3			74±3	639±68	
CONH ₂	7.5	963±80	53±1			53±1	375±29	
3.5 DMP	7.5	1098±52	58±2			58±2	579±49	

Rats fasted 18 hr before the beginning of the experiment.

TABLE 3

Treatment mg/kg os	CONTROLS				CONH ₂ 7.5 mg/kg os × 15 days			
	Blood cells				Blood cells			
	body wt. increase g	adipose tissue mg/b.w.	Red 10 ⁶ mm ³ ± S.E.	White 10 ³ mm ³ ± S.E.	body wt. increase g	adipose tissue mg/b.w.	Red 10 ³ mm ³ ± S.E.	White 10 ³ mm ³ ± S.E.
Fed	138	466	—	—	130	432	—	—
Fasted	92	386	6.42 ± 0.09	14.7 ± 1.5	96	396	5.91 ± 0.4	12.8 ± 2.1
Fasted + CONH ₂ 7.5	93	356	5.53 ± 0.42	8.8 ± 1.7	84	411	6.06 ± 0.6	15.8 ± 6

TABLE 4

Treatment mg/kg os	CONTROLS						CONH ₂ 7.5 mg/kg, os × 15 days					
	Blood cells						Blood cells					
	FFA uEq/l ± S.E.	Glucose mg/100 ml ± S.E.	TG mg/100 ml ± S.E.	CHOL mg/100 ml ± S.E.	P mg/ml ± S.E.	uEq/l ± S.E.	mg/100 ml ± S.E.	mg/100 ml ± S.E.	mg/100 ml ± S.E.	mg/ml ± S.E.	mg/ml ± S.E.	mg/ml ± S.E.
Fed	143 ± 13	100 ± 0.5	107 ± 4	50 ± 6	4.1 ± 0.05	155 ± 25	93 ± 3	109 ± 7	74 ± 7	5.2 ± 0.2	5.2 ± 0.2	5.2 ± 0.2
Fasted	288 ± 48	76 ± 6	94 ± 3	60 ± 5	4.1 ± 0.3	553 ± 70	72 ± 0.7	98 ± 3	65 ± 2	4.4 ± 0.2	4.4 ± 0.2	4.4 ± 0.2
Fasted + CONH ₂ 7.5	142 ± 25	68 ± 7	63 ± 10	61 ± 4	3.9 ± 0.4	271 ± 47	74 ± 3.9	72 ± 7	59 ± 2	3.9 ± 0.3	3.9 ± 0.3	3.9 ± 0.3

TABLE 5
Preventive effect

Treatment	Liver TG mg/100 g
Saline	504
Alcohol 2.1 ml/kg, oral	1690
Alcohol + 3.5 DMP 15 mg/kg, i.p.	1026 (°)
Alcohol + CONH ₂ 15 mg/kg, i.p.	721 (°)
Alcohol 1.6 ml/kg, oral	1161
Alcohol + CONH ₂ 15 mg/kg i.p.	552 (°)

Animals were fasted for 18 hrs. Drugs were given together with alcohol (50+ solution). Animals were killed 8 hr after the end of the treatment.

(°) P 0.01 in respect to controls.

TABLE 6
Curative effect

Treatment	Liver TG mg/100 g.
Saline	486
Alcohol 1.6 ml/kg, oral	1180
Alcohol + 3.5 DMP 15 mg/kg, i.p	1200
Alcohol + CONH ₂ 15 mg/kg, i.p.	867 (°)

Animals were fasted for 18 hrs. Drugs were given 1 hr before killing. Alcohol was given 8 hr before killing.

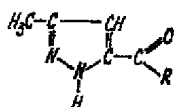
(°) P 0.01

When employed as anti-lipemic agents, the compounds of formula I are preferably administered via the oral route in the form of dragées, tablets, capsules, syrups, or elixirs. The compounds can also be administered by injection by employing a suspension of the compound in water or physiological saline, or an aqueous solution thereof or a solution of the compound concerned in a solvent consisting of aqueous propylene glycol, dimethyl guanide, sodium salicylate, methyl glucosamine or aqueous polyethylene glycol. In addition to

the active ingredients, the tablets contain fillers, extenders, lubricants and so forth of any conventional kind. Generally the active ingredient is from 25% to 90% of the total composition, by weight. Typical examples of said tablets or capsules are those containing from 20 to 200 milligrams of active ingredient, preferably 100 milligrams.

WHAT WE CLAIM IS:—

1. A process for preparing amides of 3-methyl-5-pyrazolecarboxylic acid having the general formula:



- wherein R is NH₂ or a radical of either a primary or a secondary amine, comprising the steps of treating ethyl acetopyruvate, dissolved in glacial acetic acid, with hydrazine hydrate, cooling to cause 3 - methyl - 5 - pyrazolecarboxylic acid ethyl ester to precipitate, reacting said ester with ammonia or a primary or secondary amine in stoichiometrically equivalent proportions in an aqueous alcoholic solution, in an excess of the amine, or in aqueous solution containing an excess of the amine or ammonia, refluxing for 4 to 15 hours and collecting the precipitated amide.
2. A process according to claim 1, wherein the ethyl ester of 3 - methyl - 5 - pyrazolecarboxylic acid is treated with ammonia, morpholine, piperidine, benzylamine or *para*-chlorobenzylamine.

3. A process according to claim 1 substantially as described.

4. An amide of 3 - methyl - 5 - pyrazolecarboxylic acid when produced by the process of any of claims 1 to 3.

5. The morpholide of 3 - methyl - 5 - pyrazolecarboxylic acid.

6. The piperidine of 3 - methyl - 5 - pyrazolecarboxylic acid.

7. The benzylamide of 3 - methyl - 5 - pyrazolecarboxylic acid.

8. The *para* - chlorobenzylamide of 3 - methyl - 5 - pyrazolecarboxylic acid.

9. A pharmaceutical composition comprising, in association with a pharmaceutical carrier, an amide as claimed in any one of claims 4 to 8.

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Reference has been directed in pursuance of Section 9, Subsection (1) of the Patents Act, 1949, to patent No. 1,048,104.